SPECIFICATION

A-METHOD FOR PRODUCING ELECTROLYZED WATER

FIELD OF THE INVENTION

The present invention relates to a method for producing acidic electrolyzed water and alkaline electrolyzed water by the electrolysis of water.

DESCRIPTION OF THE PRIOR ART

The production of acidic electrolyzed water and alkaline electrolyzed water by the electrolysis of the water in which chlorine electrolyte ishas been added is usually put in practicewell-known. The acidic electrolyzed water has a range of pH from 2.0 to 3.5 and has a strong sterilizing effect against a colon bacillus, various germs and bacterium so that it recently have has started to be used broadly in athe medical field, an-agricultural field, a-farming field and other fields. While, since the alkaline electrolyzed water has a range of pH from 10.5 to 12.0 and indicates strongly alkaline, it is well-known that the alkaline electrolyzed water has a weak sterilizing effect and at the same time shows strong detergency against the stain-containing oils and proteins. Therefore, the new applications of the alkaline electrolyzed water as the wash water for vegetables, fruits, dairy products and marine products, and further for mechanical parts or electronic materials—are, have gradually been developed.

To produce thesethe acidic electrolyzed water and alkaline electrolyzed water by electrolysis of water, for example, the method to-use of using an electrolyzing apparatus having a structural feature to divide of dividing a chamber tointo an anode chamber and a cathode chamber by a diaphragm and arranging an anode plate in the anode chamber and a cathode plate in the cathode chamber and to carry carrying out the electrolysis by filling the apparatus with water to which

electrolyte is has previously been added can be mentioned. Further, as the another example, thea method to use of using an electrolyzing apparatus having a structural feature to divide a chamber tointo an anode chamber, an intermediate chamber and a cathode chamber by two diaphragms and fillingintroducing high concentrated electrolyte ininto the intermediate chamber, while, fillingintroducing water ininto the anode chamber and the cathode chamber, and then carrying out $\frac{1}{2}$ can be mentioned. These methods arehave been practically used.

In these methods, the scale stuck to the cathode plate or the generation of sludge shape-precipitation in the alkaline electrolyzed water are pointed out as problems. Namely, the hardness components such as calcium or magnesium contained in water are stuckstick to the cathode plate as the scale which causes the serious problems such as the an increase of electricin the electrical resistance of the electrodes, the loading of the diaphragm or the obstruction of water flow. Up to the present time, the phenomenon thatof the scale stickssticking to the cathode ishas been considered as an unavoidable phenomenon. As a countermeasure to avoid the sticking of the scale, thea method to remove of removing the hardness components contained in the water by means of a water softener, washing the scale stuck to the cathode by an acid or to releasereleasing the scale by reversing the polarity of the electrodes has been practiced. However, the practical carrying out of these countermeasures, are not advantageous from the viewpoint viewpoints of cost or frombeing the viewpoint of the troublesome-process.

THE OBJECT OF THE INVENTION

The present invention is carried out concerning has been arrived at in view of the above-mentioned circumstances and the object of the present invention is to provide thea method for water electrolysis that can avoid the sticking of scale to the cathode and generation of sludge shape precipitation in the alkaline water atduring the production of acidic

electrolyzed water and alkaline electrolyzed water by an easy way.

DISCLOSURE OF THE INVENTION

The inventors of the present invention, have continued thean eager investigation to accomplish the above mentioned object and have found out that the sticking of scale to the cathode plate can be effectively avoided by combining two different technologies that are—strictly restricted restrict the water flow rate to the cathode chamber with regardingregard to the electric current to the cathode plate, and a water softening treatment applied only to the water which is supplied to the cathode chamber, while producing of acidic electrolyzed water in the anode chamber and alkaline electrolyzed water in the cathode chamber, and accomplished the present invention.

That is, the present invention is thea method for producing of the-electrolyzed water comprising, using an—a water-electrolyzing apparatus of water—having a structural feature to divide which divides an electrolyzer to into an anode chamber and a cathode chamber by a diaphragm and arranging an anode plate in the anode chamber and a cathode plate in the cathode chamber and to carry carrying out the electrolysis by filling the <u>electrolysis</u> with water to which electrolyte ishas previously been added, wherein the flow rate of water to be provided to the cathode chamber is restricted to 40mL(milli litermilliliter)/min. per 1A (one ampere) of loading electric current, or less, and previously softening the water provided to the cathode chamber.

Further, the present invention is thea method for producing of the electrolyzed water comprising, using an a water-electrolyzing apparatus of water having a structural feature to divide an electrolyzer to into an anode chamber, an intermediate chamber and a cathode chamber by two diaphragms and

arranging an anode plate in the anode chamber, a cathode plate in the cathode chamber and containing an electrolyte solution in the intermediate chamber,

providing water to the anode chamber and cathode chamber of said water-electrolyzing apparatus-of-water,

and generating acidic water in the anode chamber and alkaline water in the cathode chamber by loading electric current so as to make electrolysis of electrolyze the water under in the presence of electrolyte supplied by means of electrophoresis from the intermediate chamber,

wherein the flow rate of water to be provided to the cathode chamber is restricted to 40mL/min. per 1A (one ampere) of loading electric current, or less, and previously softening the water provided to the cathode chamber. Desirably, the above-mentioned water softening treatment is carried out by passing the water through the water softening apparatus in which cationic exchange resin is filled-up.

Furthermore, the present invention is thea method for producing the electrolyzed water, wherein the flow rate of the water to be provided to the anode chamber is restricted to 40mL/min. per 1A (one ampere) of loading electric current, or less. Furthermore, the present invention is thea method for producing the electrolyzed water, wherein the water for dilution is mixed towith the electrolyzed water produced in said anode chamber so-as-to prepare acidic electrolyzed water having a pH from 2.0 to 4.0 and the water for dilution is mixed towith the electrolyzed water produced in said cathode chamber so as to prepare alkaline electrolyzed water having a pH from 10 to 13.

BRIEF ILLUSTRATION OF THE DRAWINGS

Fig. 1 shows the cross sectional view of one example of the water-electrolyzing apparatus for water-used for the method of the present invention.

- Fig. 2 shows the cross-sectional view of another example of the water-electrolyzing apparatus for water used for the method of the present invention.
- Fig. 3 shows the cross-sectional view of the other another example of the water-electrolyzing apparatus for water-used for the method of the present invention.

In the drawing-each numerical, the numbers are, 1 and 2; a diaphragm, 3 and 4; an electrode plate, 5 and 9; water,

A, B and C; a wall of the electrolyzer, D; an anode chamber, E; a cathode chamber, F; an intermediate chamber, G and H; groove for water flow

DETAILED DESCRIPTION OF THE INVENTION

Fig. 1 is thea drawing showing thea cross-sectional view of one example of thea water-electrolyzing apparatus for water used for the method of the present invention. That is, Fig. 1 is the cross-sectional view of the water-electrolyzing apparatus for water providingprovided with an electrolyzer divided to into an anode chamber and a cathode chamber by one diaphragm and arrangingarranged with an anode plate tointo the anode chamber and a cathode plate tointo the cathode chamber. (A) and (B) respectively are the walls of the electrolyzer. This electrolyzer is divided to into the anode chamber (D) and the cathode chamber (E). (3) and (4) is are the electrode $\frac{\text{plate}}{\text{plate}}$ plates, and electrode (3) is used as an anode plate, and electrode (4) is used as a cathode plate. And 6', 7', $10'_{\tau}$ and 11' are valves for adjusting the flow laterate of the water.

The method for producing of anionic electrolyzed water and cationic electrolyzed water produced-by the electrolytic treatment of water will be illustrated using the water electrolyzing apparatus providing with the electrolyzer of Fig. 1. The water (5) provided to an anode side is divided to theinto water to be electrolyzed (6) and the water not to be electrolyzed (7). To the water to be electrolyzed (6), a small amount of electrolyte is added and introduced to the anode

chamber (D). This water to be electrolyzed (6) is electrolyzed in the anode chamber (D) and becomes the acidic electrolyzed water. The obtained acidic electrolyzed water can be used as is, or can be joined with the water not to be electrolyzed (7) and diluted to the desired pH (for example, pH2.0-4.0) so that to be used as the acidic electrolyzed water (8). In the meanwhile, the water provided to the cathode side (9) is divided to into the water to be electrolyzed (10) and the water not to be electrolyzed (11). To the water to be electrolyzed (10), a small amount of electrolyte is added and provided to the anode chamber (E). This water to be electrolyzed (10) is electrolyzed in the cathode chamber and becomes alkaline electrolyzed water. The obtained alkaline electrolyzed water can be used as is, or can be joined with the water not to be electrolyzed (11) and diluted to the desired pH (for example, pH10.0-13.0) so that to be used as the alkaline electrolyzed water (12).

Fig._2 shows the cross-sectional view of thea water electrolyzing apparatus providing with thean electrolyzer arranging having arranged therein an anode chamber and a cathode chamber separated by dividing with two diaphragms, used in the present invention. (A), (B) and (C) are respectively indicating a wall of the electrolyzer. This electrolyzer is divided tointo an anode chamber (D), an intermediate chamber (F) and a cathode chamber (E) by two diaphragms (1) and (2). While (3) and (4) are an-electrode plateplates, and the electrode plate (3) is the anode plate and the electrode plate (4) is the cathode plate. The electrode plate (3) and the diaphragmsdiaphragm (1), and the electrode plate (4) and the diaphragmsdiaphragm (2) can be separated or can be contacted. Fig. 2 is the case that the electrode plateplates and the diaphragms are tightly contacted. As mentioned later, in the case when the electrode plateplates and the diaphragms are contacted, it is desirable to use an electrode plate that has holes and an electrically non-conductive material is put in between said electrode

plates and said diaphragms or to use an electrode whose face is coated by an electrically non-conductive material. In the intermediate chamber (F), electrolyte aqueous solution of high concentration is filled up. Ordinary Ordinarily, an aqueous solution of sodium chloride or potassium chloride of over than 10% concentration is used. Further, said aqueous solution can be provided to into the intermediate chamber (F) from the separated tank for electrolyte aqueous solution using a pump. The concentration of the electrolyte can be higher in with the limit being not to obstruct the fluidity of the aqueous solution. Further, 6', 7', 10' and 11' are the valves to adjust the amount of individual water flows to the anode and cathode chamberchambers and diluting water of acid and alkaline products.

The method for producing acidic electrolyzed water and alkaline electrolyzed water by electrolyzing of water will be illustrated more in detail according to the electrolyzing apparatus of water providing withutilizing the electrolyzer of Fig. 2. The water (5) provided to the anode side is divided totwo-into water that are to be electrolyzed (6) and not to be electrolyzed (7). The water to be electrolyzed (6) is led to the anode chamber (D). To the water to be electrolyzed (6), electrolyte is provided by electrophoresis from the intermediate chamber (F), and the water is electrolyzed and becomes acidic electrolyzed water. The obtained acidic electrolyzed water can be used as is, or can be joined with the water not to be electrolyzed (7) and diluted to the desired pH (for example, pH2.0-4.0) so that to tan be used as the acidic electrolyzed water (8). In the meanwhile, the water provided to the cathode side (9) is divided to the water to be electrolyzed (10) and the water not to be electrolyzed (11). The water to be electrolyzed (10) is led to the cathode chamber (E). To the water to be electrolyzed (10), electrolyte is provided by electrophoresis from the intermediate chamber (F), and the water is electrolyzed and becomes alkaline electrolyzed water. The obtained alkaline

electrolyzed water can be used as is, or can be joined with the water not to be electrolyzed (11) and diluted to the desired pH (for example, pH10.0-13.0) so that toit can be used as the alkaline electrolyzed water (12).

In the present invention, atduring the electrolysis of water by means of the electrolyzing apparatus of water shown in Fig. 1 or Fig. 2, the water to be provided to the cathode chamber (E) is necessary has to satisfy following two points. That is, the first one is to restrict the flow rate of the water to be provided to the anode chamber to 40mL/min. per 1A of loading electric current or less. And the second one is that the water to be provided to the cathode chamber is the water that ishas been softened previously. By satisfying said two points, the sticking of scale to the cathode can be effectively avoided and the generation of sludge shape precipitation in the alkaline electrolyzed water can be prevented, further, the blockade trouble by precipitation of a pipe or a tank can be prevented.

Further, in the present invention, regarding the water to be provided to the cathode chamber (E), it is desirable to restrict the providing rate of water to 40mL/min. per 1A (ampere) of loading electric current or less and it is also desirable to restrict the providing rate of water to the anode chamber (D) to 40mL/min. per 1A (ampere) of loading electric current or less. By restricting the amount of water for electrolysis as above, the transfusing phenomenon of water from the anode to the cathode occurs at occurring during the electrolysis can be prevented and can elevate the concentration of free chlorine contained in the acidic electrolyzed water.

Above The above-mentioned water softening treatment can be conveniently carried out by passing the water through a water softening apparatus in which a cationic exchange resin is filled-up. For the softening treatment of the water to be provided in the cathode chamber (E) (water 10 to be electrolyzed), it is preferable to arrange a water softening

apparatus in which a cationic exchange resin is filled-up, located in between the valve (10') and the cathode chamber (E) of the electrolyzer. As a cationic exchange resin, thea cationic exchange resin using a copolymer consisting of styrene and divinylbenzene or a copolymer consisting of methacrylic acid and divinylbenzene as a mother resin and introducing an acidic group such as a sulfone group or carboxylic group to said mother resin as the exchanging group is used.

The object of the present invention can be accomplished by softening the water to be provided to the cathode chamber alone, among the whole water to be provided to the electrolyzing apparatus of water. The water to be provided to the cathode chamber to be electrolyzed (10) alone, which is a part of the water to be provided to the electrolyzing apparatus of water, is previously softened by passing the water through a water softening apparatus. And in In the present invention, since the providing amount of water to be electrolyzed (10) is 40mL/min. per 1A of loading electric current or less and is recognized to be small, the size of athe water softening apparatus in which the cationic exchange resin is filled up—can be minimized and the cycle for the washing of the cationic exchange resin can be extended. Therefore, by the present invention, the above-mentioned operations are coupled together and effectively prevent the sticking of scale to the cathode.

For example, to generate 1000mL of acidic electrolyzed water and alkaline electrolyzed water respectively every minute, it is necessary to provide 2000mL of water to an electrolyzer every minute, therefore, Therefore, in the case to soften whole of softening all the water to be provided to the electrolyzer, it is necessary to soften 2000mL of water every minute. While, in the case of the present invention, the flow rate of water to be provided to the cathode and to be electrolyzed is 40mL/min. per 1A of loading electric current or less. This rate is converted to the case that produces

1000mL of acidic electrolyzed water and alkaline electrolyzed water respectively every minute. Since the electric current value loaded at the electrolysis process is generally approximately 6-10 ampereamperes, the amount of water to be provided to the cathode and to be electrolyzed is 240mL or less in the case of 6 ampereamperes and 400mL or less in the case of 10 ampereamperes. That is, in the case of the present invention, the maximum amount of water for softening is 400mL per minute. This amount is less than 1/5 to 2000mL/min that is the necessary amount for softening of the conventional type. Therefore, in the present invention, the minimization of the size of a water softening apparatus becomes possible and the cycle for the washing of the cationic exchange resin can be extended.

The electrode and the diaphragm of the electrolyzing apparatus of water used in the present invention will be illustrated. The electrode and the diaphragm can be contacted or can be not contacted. In the case when the electrode and the diaphragm are used in contacted condition, a plate having various holes or a net is desirably used as an electrode. In the case when the electrode and the diaphragm are used with a distance therebetween, it is not necessary to have a hole. As the material of the electrode, for example, a plate of copper, lead, nickel, chrome, titanium, tantalum, gold, platinum, iron oxide, stainless steel, carbon fiber or graphite can be mentioned, in particular, as the material of the anode, a platinum genus metal-plated or baked titanium is desirably used. Further, as the material of the cathode, platinum-plated titanium is desirably used, however, chrome stainless steel (SUS316L) or nickel can be also used.

Still further, when the above-mentioned electrode plate with various holes is used in contact with a diaphragm, it is desirable to use an electrode plate prepared by arranging a sheet shape non-electrically conductive material which has corresponding holes to the electrode plate between each electrode plate and diaphragm, or to use an electrode plate

with many holes to the surface of which facing to a faces the diaphragm isbeing coated by a non-conductive film. As the concrete examplespecific examples of the material used for the sheet shape non-electrically conductive material, are synthetic resinresins such as a fluororesin (registered Trade Mark: Teflon), ABS resin, acrylic resin, epoxy resin, polyurethane resin, polypropylene resin, nylon resin, polyethyleneterephthalate resin, polyamide resin and vinyl chloride resin, or natural rubber or elastomer such as SBR, chloroprene and polybutadiene. These electrode plates are disclosed in Japanese Patent Laid open publication 8-276184. These types of electrode plates are desirable to use because they do not generate electrolysis of water at the surface of the diaphragm side, therefore, the phenomenon that the staying of the gases staying between the electrode and diaphragm and abstracting obstruction of the flow of electric current can be reduced.

As the diaphragm, thea material that has water permeability can be used, for examples, woven cloth or nonwoven cloth such as polyvinylfluoride fiber, asbestos, glass wool, polyvinylchloride fiber, polyvinylidenechloride fiber, polyester fiber or aromatic polyamide fiber. As the another examples example, which forms the diaphragm withby mixing of woven cloth, non-woven cloth of polyester fiber, nylon fiber or polyethylene fiber as an aggregate, and chlorinated polyethylene, polyvinylchloride or polyvinylidenechloride are used as a film, or a diaphragm prepared with mixing of titanium oxide to said diaphragm can be mentioned. Furthermore, a semi-permeable membrane such as cellophane, cationic ion-exchange membranemembranes or anion-exchange membrane membranes can be used. The electrolysis condition of the present invention is to charge a high load electric current to the small amount of water to be electrolysis so as to generate very strong acidic or alkaline water and generate highly concentrated chlorine gas, it is desirable to

select a diaphragm that can endure to the severe conditions.

In the electrolyzing apparatus of water of Fig. 2, as shown in Fig. 3, the edge part of the anode chamber (D) is partitioned by a partition board 13 so as to form a groove (G), and the edge part of the cathode chamber (E) is partitioned by a partition board 14 so as to form a groove (H). Water not to be electrolyzed (7) can be flownflow in the groove (G) and water not to be electrolyzed (11) can be flownflow in the groove (H), and the water that flows in groove (G) and groove (H) acts conveniently as the coolant of the electrolyzer.

EXAMPLES

Example 1

The Example which uses the electrolyzing apparatus of water of Fig. 2 will be substantially illustrated as follows. The size of the electrolyzer is; 15cm in length-direction, 9cm in width-direction and 6cm in thickness. As the electrode plate for anode (3), a platinum/iridium oxide baked titanium plate having many holes and whose actual surface area is $50\,\mathrm{cm}^2$ is used. While, as the electrode plate for cathode (4), a platinum-plated titanium plate having many holes and whose actual surface area is 50cm^2 is used. AtDuring the actual use, a sheet of fluororesin (registered Trade Mark: Teflon), which is an electrically non-conductive material, having many holes is coated by a non-conductive film toonto the diaphragm side of each electrode plate. As the diaphragm (1) used for the partition of the anode chamber (D) and the intermediate chamber (F), an anionic ion-exchange membrane is used and as the diaphragm (2) used for the partition of the cathode chamber (E) and the intermediate chamber (F), a cationic ionexchange membrane is used. In the intermediate chamber (F), an aqueous solution of approximately 30% sodium chloride is filled up as the electrolyte.

As the water (5) to be provided to the anode, city water is used, and is divided to theinto water to be electrolyzed (6) and the water not to be electrolyzed (7). The water to be electrolyzed (6) is introduced to the anode chamber (D) and generates acidic electrolyzed water by electrolysis. The obtained acidic electrolyzed water is joined and mixed with not electrolyzed water (7) and adjusted to the desired pH and flownflows out from the outlet (8), thus the acidic electrolyzed water of a desired pH is obtained. Further, as the water (9) to be provided to the cathode, city water is used, and is divided to the water to be electrolyzed (10) and the water not to be electrolyzed (11). And the The water to be electrolyzed (10) alone is softened by passing through a softening apparatus in which a cationic exchange resin is filled up—and introduced to the cathode chamber (E) and electrolyzed. Thus the alkaline electrolyzed water is generated. This alkaline electrolyzed water is joined with not electrolyzed water (11) and adjusted to the desired pH and flownflows out from the outlet (12), thus the alkaline electrolyzed water of a desired pH is obtained.

The direct electric current loaded to the electrode is set to 6.5 ampereamperes and the voltage at the operation is 6.7volt volts. The flow rate of water (6) to be electrolyzed and to be introduced to the anode chamber is adjusted to 100mL per minute, further, the flow rate of water (7) not to be electrolyzed is adjusted to 900mL per minute, and they are then joined and mixed together with at the outlet and 1000mL per minute of acidic electrolyzed water is obtained. The pH value of the obtained acidic electrolyzed is 2.68, the ORP value is 1130mV and the measured value of contained free chlorine is 30ppm. While, the flow rate of water (10) to be electrolyzed and to be introduced to the cathode chamber is adjusted to 100mL per minute, further, the flow rate of water (11) not to be electrolyzed is adjusted to 900mL per minute, and they are then joined and mixed together with at the outlet and 1000mL per minute of alkaline electrolyzed water is

obtained. The pH value of the obtained alkaline electrolyzed water is 11.54. Maintaining the same operating condition, the electrolysis experiment is continuously carried out for 48 hours, and the sticking of scale to the cathode is not observed at all. Further, the generation of precipitation is not observed in the obtained alkaline electrolyzed water.

Comparative Example 1

In Example 1, the flow rate of water to be introduced to the cathode chamber (10) is adjusted to 100mL per minute, while in Comparative Example 1, the flow rate of water to be introduced to the cathode chamber (10) is adjusted to 1000mL per minute and the flow rate of water (11) not to be electrolyzed is adjusted to OmL per minute. Other conditions are set the same as tointo Example 1 and the electrolyzed water is produced.

After starting the electrolysis experiment, the voltage starts to elevate along with the time lapse, and after 48hours it becomes impossible to continue the electrolysis experiment because of high voltage. The reason whyfor the phenomena can be considered is to be as follows. That is, because the hardness component is remaining remains in the water (10) to be introduced to into the cathode chamber (E), and the remaining hardness component stucksticks to the cathode plate as the scale.

Comparative Example 2

In Example 1, the water to be electrolyzed (10) is introduced to the cathode chamber (E) after passing through a softening apparatus in which cationic exchange resin is filled up-and the water is softened, while in the Comparative Example 2, the water to be electrolyzed (10) is introduced to into the cathode chamber (E) without softening. Other conditions are set the same as tointo Example 1 and the electrolyzed water is produced.

5000mL specimen of the alkaline electrolyzed water is picked out respectively from the alkaline electrolyzed water produced in Example 1 and Comparative Example 2. Each specimen is filtrated using two filtering paperpapers (product of Tokyo Roshi Co., Ltd., Trade Mark "advantec") and weighted after dried updrying so that the residue can be measured. The results are summarized in Table 1.

Table 1

	weight of	weight of	weight change	required
	filter paper	filter paper	before and	time to
	before	after	after	filter
	filtration	filtration	filtration	5000mL
	(g)	(g)	(g)	
Example 1	1.6529	1.7183	0.0654	50 minutes
Co. Example 2	1.6238	2.0154	0.3923	6 hours

It is clearly understood from Table 1 that the amount of precipitate in the alkaline electrolyzed water produced in Comparative Example 2 was greater than the amount of precipitate in the alkaline electrolyzed water produced in Example 1. And the The filtering time for the alkaline electrolyzed water produced in Comparative Example 2 iswas remarkably longer than for Example 2, because the filter paper of Example 2 iswas plugged with the precipitate. After the filtration, thea yellowish adhesion is was observed on the filter paper. Further, according to the results of Table 1, the amount of scale contained in the alkaline electrolyzed water of the examples iswas calculated. The amount of scale contained in the alkaline electrolyzed water produced in Example 1 iswas 13 ppm and that of Comparative Example 2 iswas 78 ppm.

EFFECT OF THE INVENTION

In the case of conventional methodmethods for producing electrolyzed water; there are problems of the sticking of scale to a cathode plate atduring the electrolysis operation and the sludge shape precipitation in the alkaline

electrolyzed water. However, according to the present invention, the sticking of scale to the cathode plate and the generation of the sludge shape precipitation in the alkaline electrolyzed water can be effectively avoided, and ean effectively produce the acidic electrolyzed water and alkaline electrolyzed water can be effectively produced.